Amendments To The Claims

1-59. (Canceled)

- 60. (Previously presented) A method for treating a B cell leukemia comprising administering a therapeutically effective amount of:
 - (i) an anti CD20 antibody having B-cell depleting activity, or a fragment thereof having B-cell depleting activity; and
 - (ii) an anti CD40L antibody that antagonizes interaction of CD40 and CD40L, or a fragment thereof that antagonizes interaction of CD40 and CD40L.

61-64. (Canceled)

- 65. (Previously presented) The method of claim 60 wherein said B cell leukemia is selected from the group consisting of chronic B cell leukemia, acute lymphoblastic leukemia, chronic lymphocytic leukemia, and Burkitt's type leukemia.
- 66. (Previously presented) The method of claim 60 wherein said anti CD20 antibody is a humanized, primatized, or chimeric anti CD20 antibody.
- 67. (Previously presented) The method of claim 60 wherein said anti CD40L antibody is a humanized, primatized, or chimeric anti CD40L antibody.
- 68. (Previously presented) The method of claim 67 wherein said anti CD40L antibody is humanized MAb 89-76 or humanized MAb 24-31.
- 69. (Previously presented) The method of claim 60 wherein said anti-CD20 antibody is radiolabeled with 90Y.

70. (Canceled)

- 71. (Previously presented) The method of claim 60, which further comprises administering at least one alkylating chemotherapeutic agent.
- 72. (Previously presented) The method of claim 71 wherein said alkylating chemotherapeutic agent is selected from the group consisting of cyclophosphamide, chlorambicil, procarbazine, and lomustine.

73-79. (Canceled)

- 80. (Previously presented) The method of claim 60 wherein the anti CD20 antibody is administered prior to the anti CD40L antibody.
- 81. (Previously presented) The method of claim 60 wherein the anti CD20 antibody is administered after the anti CD40L antibody.
- 82. (Previously presented) The method of claim 80 wherein the anti CD20 antibody is IDEC-C2B8 and the anti CD40L antibody is humanized MAb 24-31.
- 83. (Previously presented) The method of claim 81 wherein the anti CD20 antibody is IDEC-C2B8 and the anti CD40L antibody is humanized MAb 24-31.
- 84. (Previously presented) The method of claim 80 which further comprises administering at last one alkylating chemotherapeutic agent.
- 85. (Previously presented) The method of claim 81 which further comprises administering at last one alkylating chemotherapeutic agent.
- 86. (Previously presented) The method of claim 66 wherein said anti CD20 antibody is IDEC-C2B8.
- 87. (Previously presented) The method of claim 86 wherein said anti CD40L antibody is humanized MAb 24-31.

- 88. (Previously presented) The method of claim 68 wherein said anti CD40L antibody is humanized MAb 24-31.
- 89. (Previously presented) The method of claim 60, wherein a weekly dose of said anti CD20 antibody is 0.4 to 20 mg/kg body weight, and a dose of said anti CD40L antibody is 0.5 to 10 mg/kg body weight.
- 90. (Previously presented) The method of claim 69 wherein said anti-CD20 antibody that is radiolabeled with 90Y is a non-chimeric murine antibody.
- 91. (Previously presented) The method of claim 84 wherein said alkylating chemotherapeutic agent is selected from the group consisting of cyclophosphamide, chlorambicil, procarbazine, and lomustine.
- 92. (Previously presented) The method of claim 85 wherein said alkylating chemotherapeutic agent is selected from the group consisting of cyclophosphamide, chlorambicil, procarbazine, and lomustine.
- 93. (Previously presented) A method for treating a B cell leukemia comprising administering a therapeutically effective amount of:
 - (i) IDEC-C2B8, and
 - (ii) humanized MAb 24-31.
- 94. (Previously presented) The method of claim 93 wherein IDEC-C2B8 is administered prior to humanized MAb 24-31.
- 95. (Previously presented) The method of claim 93, wherein a weekly dose of said IDEC-C2B8 is 0.4 to 20 mg/kg body weight, and a dose of said humanized MAb 24-31 is 0.5 to 10 mg/kg body weight.
- 96. (Previously presented) The method of claim 93 wherein said B cell leukemia is selected from the group consisting of chronic B cell leukemia, acute lymphoblastic leukemia, chronic lymphocytic leukemia, and Burkitt's type leukemia.

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- 97. (Previously presented) The method of claim 93 wherein IDEC-C2B8 is radiolabeled with 90Y.
- 98. (Previously presented) The method of claim 93 which further comprises administering at least one alkylating chemotherapeutic agent.
- 99. (Previously presented) The method of claim 98 wherein said alkylating chemotherapeutic agent is selected from the group consisting of cyclophosphamide, chlorambicil, procarbazine, and lomustine.